

be maintained either as individual records or, in food animal practices, on a group, herd, flock, or per-client basis. Records shall be adequate to provide the following information:

- (1) The established name of the drug and its active ingredient, or if formulated from more than one ingredient, the established name of each ingredient;
- (2) The condition treated;
- (3) The species of the treated animal(s);
- (4) The dosage administered;
- (5) The duration of treatment;
- (6) The numbers of animals treated; and
- (7) The specified withdrawal, withholding, or discard time(s), if applicable, for meat, milk, eggs, or any food which might be derived from any food animals treated.

(b) A veterinarian shall keep all required records for 2 years or as otherwise required by Federal or State law, whichever is greater.

(c) Any person who is in charge, control, or custody of such records shall, upon request of a person designated by FDA, permit such person designated by FDA to, at all reasonable times, have access to, permit copying, and verify such records.

#### **Subpart B—Rules and Provisions for Extralabel Uses of Drugs in Animals**

##### **§ 530.10 Provision permitting extralabel use of animal drugs.**

An approved new animal drug or human drug intended to be used for an extralabel purpose in an animal is not unsafe under section 512 of the act and is exempt from the labeling requirements of section 502(f) of the act if such use is:

- (a) By or on the lawful written or oral order of a licensed veterinarian within the context of a valid veterinarian-client-patient relationship; and
- (b) In compliance with this part.

##### **§ 530.11 Limitations.**

In addition to uses which do not comply with the provision set forth in § 530.10, the following specific extralabel uses are not permitted and

result in the drug being deemed unsafe within the meaning of section 512 of the act:

- (a) Extralabel use in an animal of an approved new animal drug or human drug by a lay person (except when under the supervision of a licensed veterinarian);
- (b) Extralabel use of an approved new animal drug or human drug in or on an animal feed;
- (c) Extralabel use resulting in any residue which may present a risk to the public health; and
- (d) Extralabel use resulting in any residue above an established safe level, safe concentration or tolerance.

##### **§ 530.12 Labeling.**

Any human or animal drug prescribed and dispensed for extralabel use by a veterinarian or dispensed by a pharmacist on the order of a veterinarian shall bear or be accompanied by labeling information adequate to assure the safe and proper use of the product. Such information shall include the following:

- (a) The name and address of the prescribing veterinarian. If the drug is dispensed by a pharmacy on the order of a veterinarian, the labeling shall include the name of the prescribing veterinarian and the name and address of the dispensing pharmacy, and may include the address of the prescribing veterinarian;
- (b) The established name of the drug or, if formulated from more than one active ingredient, the established name of each ingredient;
- (c) Any directions for use specified by the veterinarian, including the class/species or identification of the animal or herd, flock, pen, lot, or other group of animals being treated, in which the drug is intended to be used; the dosage, frequency, and route of administration; and the duration of therapy;
- (d) Any cautionary statements; and
- (e) The veterinarian's specified withdrawal, withholding, or discard time for meat, milk, eggs, or any other food which might be derived from the treated animal or animals.